## AMENDMENTS TO THE CLAIMS

- 1-4. (Canceled).
- 5. (Currently Amended) A method of inducing a prophylactically effective an immune response against a *Helicobacter pylori* polypeptide or inducing an immune response for reducing the degree of *Helicobacter pylori* infection in a primate mammal, said method consisting essentially of administering to said primate mammal an a prophylactically effective amount of a prophylactically effective *Helicobacter pylori* polypeptide antigen by the subdiaphragmatic, strict systemic route.
- 6. (Previously Presented) The method of Claim 5, in which a Th1-type immune response is induced by said subdiaphragmatic, systemic administration.
- 7. (Currently Amended) The method of Claim 6, <u>further comprising induction of wherein</u> a Th1-type immune response and a Th2-type immune response, <u>wherein are induced and</u> the immune response of said <u>primate mammal</u> is characterized by either (i) a ratio of the ELISA IgG2a:IgG1 titers greater than or equal to 1:100, or (ii) a ratio of the ELISA IgG2a:IgA titers greater than or equal to 1:100.
- 8. (Previously Presented) The method of Claim 7, in which the immune response of said primate mammal is characterized either (i) by a ratio of the ELISA IgG2a:IgG1 titers greater than or equal to 1:10, or (ii) by a ratio of the ELISA IgG2a:IgA titers greater than or equal to 1:10.

- 9. (Previously Presented) The method of Claim 8, in which the immune response of said primate mammal is characterized either (i) by a ratio of the ELISA IgG2a:IgG1 titers greater than or equal to 1:2, or (ii) by a ratio of the ELISA IgG2a:IgA titers greater than or equal to 1:2.
  - 10. (Canceled).
- 11. (Previously Presented) The method of Claim 10, in which the *Helicobacter pylori* antigen comprises the UreB or UreA subunit of a *Helicobacter pylori* urease.
  - 12-14. (Canceled).
- 15. (Currently Amended) The method of Claim 5, in which <u>said</u> [[the]] *Helicobacter pylori* <u>polypeptide</u> antigen is administered by a <u>strict</u> systemic route selected from the subcutaneous route, the intramuscular route, and the intradermal route.

16 and 17. (Canceled).

- 18. (Currently Amended) The method of Claim 5, in which <u>said</u> [[the]] *Helicobacter* pylori polypeptide antigen is administered in the dorsolumbar region of said <u>primate</u> mammal.
  - 19-24. (Canceled).

25. (Previously Presented) A method of inducing a prophylactically effective an immune response against a Helicobacter polypeptide comprising the UreB or UreA subunit of a Helicobacter pylori urease or inducing an immune response for reducing the degree Helicobacter infection in a mammal, said method comprising in order the steps of:

mucosally administering, in an initial immunization, <u>an a prophylaetically</u> effective effective amount of <u>a Helicobacter pylori</u> polypeptide comprising the UreB or UreA subunit of a <u>Helicobacter pylori</u> urease a prophylaetically effective <u>Helicobacter pylori</u> polypeptide antigen to said mammal to prime an immune response; and then

parenterally administering <u>an</u> a <u>prophylactically</u> effective amount of <u>a Helicobacter pylori</u> <u>polypeptide comprising the UreB or UreA subunit of a Helicobacter pylori urease</u> a <u>prophylactically effective Helicobacter pylori polypeptide antigen</u> to said mammal to boost said immune response.

26-36. (Canceled).

- 37. (Previously Presented) The method of claim 25, further comprising carrying out more than one mucosal administration.
- 38. (Previously Presented) The method of claim 25, further comprising carrying out more than one parenteral administration.

39. (Canceled).

- 40. (Previously Presented) The method of Claim 25, in which the mucosal administration is oral administration.
  - 41-44. (Canceled).
- 45. (Currently Amended) The method of Claim 25, further comprising mucosally coadministering a mucosal adjuvant selected from the group consisting of *Escherichia coli* heat labile enterotoxin (LT), cholera toxin (CT), *Clostridium difficile* toxin, *Pertussis* toxin (PT), and combinations, subunits, toxoids, and mutants derived therefrom with <u>said</u> [[the]] mucosally administered *Helicobacter pylori* polypeptide comprising the UreB or UreA subunit of a *Helicobacter pylori* urease antigen.
- 46. (Currently Amended) The method of Claim 25, in which a parenteral adjuvant selected from the group consisting of alum, QS-21 (purified fraction of saponin extracted from *Quillarja Saponaria Molina*), DC-CHOL (3-beta-(N-(N',N'-dimethylamino-ethane)carbamoyl)cholesterol), and BAY R1005 (N-(2-deoxy-2-L-leucylamino-beta-D-glucopyranosyl)-N-octa-decyldodecanoylamide acetate) is co-administered with <u>said</u> [[the]] parenterally administered *Helicobacter pylori* polypeptide comprising the UreB or UreA subunit of a *Helicobacter pylori* urease antigen.
- 47. (Previously Presented) The method of Claim 25, in which the parenteral administration is intramuscular administration or subcutaneous administration.

48. (New) The method of claim 5, wherein said primate is a human.